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assay gene function in progenitor cells of the selected lineage, and/or (ii) to render selected cells more suitable for transplantation.

- 5. (Amended) A method according to <u>Claim</u> [any of Claims] 1 [to 4] further comprising:-
 - (iv) introducing into the multipotential cell a second selectable marker that is differentially expressed in cells of a selected sub-lineage compared with its expression in other cells, wherein cells of the selected sub-lineage are formed by differentiation of cells of the selected progenitor lineage; and
 - (v) when a culture of progenitor cells of the selected lineage has been obtained, allowing or inducing differentiation of the cells and selecting for cells that express the second selectable marker.
- 6. (Amended) A method according to <u>Claim</u> [any of Claims] 1 [to 5] wherein the selectable marker is introduced into the multipotential cell by targeted integration or random gene trap integration so as to be operatively coupled to a gene that is differentially expressed in progenitor cells of the selected lineage.
- 7. (Amended) A method according to <u>Claim</u> [any of Claims] 1 [to 5] wherein the selectable marker is introduced into the multipotential cell via random integration of a transgene in which the selectable marker is operatively coupled to a gene that is differentially expressed in progenitor cells of the selected lineage.
- 8. (Amended) A method according to <u>Claim</u> [any of Claims] 1 [to 7] wherein the multipotential cell is an ES, EG or EC cell and the method comprises forming an embryoid body, or otherwise inducing differentiation of the cells.
- 10. (Amended) A method according to Claim 8 [or 9] wherein differentiated cells of an embryoid body are dissociated using a protease, such as trypsin.
- 11. (Amended) A method according to <u>Claim</u> [any of Claims] 1 [to 10], for generating a culture that is purified or enriched in respect of neural progenitors,

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comprising introducing into the multipotential cell a selectable marker that is differentially expressed in neural progenitor cells.

- 14. (Amended) A method according to <u>Claim</u> [any of Claims] 1 [to 10] for generation of cardiac progenitor cells, wherein the selectable marker is expressed in cells that express the Nkx 2.5 or GATA-4 gene.
- 15. (Amended) A method according to <u>Claim</u> [any of Claims] 1 [to 10] for generating a culture that is purified on enriched in respect of haematopoietic progenitors.
- 17. (Amended) A method according to <u>Claim</u> [any of Claims] 1 [to 16] wherein the selectable marker is an antibiotic resistance gene and the method comprises culture in the presence of antibiotic.
- 22. (Amended) A method according to Claim 20 [or 21] to assay whether the factor has a proliferative, maturation, toxic or protective effect on progenitor cells of the selected lineage.
- 24. (Amended) A method of preparing a neural progenitor cell or a differentiated progeny thereof for storage, comprising obtaining the cell in a method according to Claim 11 [any of Claims 1 to 17] and freezing the cell in the presence of a cryoprotectant.
- 25. (Amended) A method of generating purified neurons, comprising obtaining a culture purified in respect of neural progenitors, using the method of <u>Claim</u> [any of Claims] 1 [to 17] wherein the selectable marker is differentially expressed in cells that express a Sox gene, and culturing the progenitors obtained in the presence of medium suitable for differentiation of the progenitor into neurons.

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